AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

- 1-30. (Canceled).
- 31. (Currently Amended) An inhibitor of HIV replication, comprising an antiviral peptide, wherein:

the antiviral peptide consists of or analog comprising a decapeptide, saida decapeptide containing (from the N-terminus to the C-terminus) a basic amino acid in at position 1—; an acidic amino acid in at positions 2 and 5; and a tryptophan in at positions 4, 7, and 8; wherein:

the amino acid at position 3 is a threonine, isoleucine or valine at position 3; a the amino acid at position 6 is threonine, alanine, and or glutamine at position 6; a the amino acid at position 9 is threonine, alanine, valine, isoleucine, methionine, or aspartate at position 9; and the amino acid at position 10 is glutamate, aspartate or asparagine at position 10; and

the decapeptide inhibits the dimerization of HIV reverse transcriptase.

- 32. (Currently Amended) The inhibitor of claim 31, wherein the basic amino acid in at position 1 is lysine or arginine.
- 33. (Currently Amended) The inhibitor of claim 31, wherein the acidic amino acid in at position 2 is glutamate.
- 34. (Previously Presented) The inhibitor of claim 31, wherein the amino acid at position 5 is glutamate.

- 35. (Currently Amended) A pharmaceutical composition comprising the The inhibitor of HIV replication of claim 31, further comprising and a pharmaceutically acceptable excipient.
- 36. (New) The inhibitor of claim 31, further comprising a vector that allows penetration of the antiviral peptide into a mammalian cell.
- 37. (New) The inhibitor of claim 36, wherein the vector is selected from the group consisting of: a liposome, a polymeric protein-binding cation, a protein, a peptide, a microparticle, and a nonoparticle.
 - 38. (New) The inhibitor of claim 37, wherein the vector is a peptide.
- 39. (New) The inhibitor of claim 38, wherein the peptide is an MPG peptidyl carrier.
- 40. (New) The inhibitor of claim 39, wherein the MPG peptidyl carrier comprises SEQ ID NO: 2 or SEQ ID NO: 3.
- 41. (New) The inhibitor of claim 39, wherein the MPG peptidyl carrier and the antiviral peptide are in the form of a complex.
- 42. (New) The inhibitor of claim 41, wherein the complex comprises the MPG peptidyl carrier and the antiviral peptide at a ratio of about 20 molecules of the MPG peptidyl carrier for 1 molecule of the antiviral peptide.
- 43. (New) The inhibitor of claim 31, wherein the antiviral peptide is SEQ ID NO: 1.
- 44. (New) The inhibitor of claim 42, wherein the antiviral peptide is SEQ ID NO: 1.

- 45. (New) An inhibitor of HIV replication comprising a chimeric peptide, wherein the chimeric peptide comprises:
- (a) a decapeptide containing (from the N-terminus to the C-terminus) a basic amino acid at position 1; an acidic amino acid at positions 2 and 5; a tryptophan at positions 4, 7, and 8; a threonine, isoleucine or valine at position 3; a threonine, alanine, or glutamine at position 6; a threonine, alanine, valine, isoleucine, methionine, or aspartate at position 9; and a glutamate, aspartate or asparagine at position 10; wherein the decapeptide inhibits the dimerization of HIV reverse transcriptase, and
 - (b) an MPG peptidyl carrier peptide.
- 46. (New) The inhibitor of claim 45, wherein the basic amino acid in at position 1 is lysine or arginine.
- 47. (New) The inhibitor of claim 45, wherein the acidic amino acid in at position 2 is glutamate.
- 48. (New) The inhibitor of claim 45, wherein the amino acid at position 5 is glutamate.
- 49. (New) The inhibitor of claim 45, further comprising a pharmaceutically acceptable excipient.
- 50. (New) The inhibitor of claim 45, wherein the MPG peptidyl carrier peptide is SEQ ID NO: 2 or SEQ ID NO: 3.
- 51. (New) The inhibitor of claim 45, wherein the decapeptide is SEQ ID NO: 1.
- 52. (New) The inhibitor of claim 45, wherein the chimeric peptide is SEQ ID NO: 4.

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53. (New) The inhibitor of claim 45, wherein the chimeric peptide is SEQ ID NO: 6.